Amendments to the Claims

Claims 1-50 (Canceled)

- 51. (Currently Amended) A chimeric or humanized anti-α5β1 integrin antibody comprising: a heavy chain variable region comprising an amino acid sequence having at least 95% identity to a sequence selected from the group consisting of SEQ ID NOs: 1, and 16; a light chain variable region comprising an amino acid sequence having at least 95% identity to a sequence selected from the group consisting of SEQ ID NOs: 7, and 18; and a constant region, wherein the source of the constant region is a human IgG.
- 52. (Previously Presented) The chimeric or humanized anti-α5β1 integrin antibody of claim 51, wherein the source of the constant region is human IgG4 or IgG2M3.
- 53. (Previously Presented) The chimeric or humanized anti-α5β1 integrin antibody of claim 52, wherein the source of the constant region is human IgG4.
- 54. (Previously Presented) The chimeric or humanized anti-α5β1 integrin antibody of claim 52, wherein the heavy chain variable region sequence comprises SEQ ID NO: 1, and the light chain variable region sequence comprises SEQ ID NO: 7.
- 55. (Currently Amended) An anti-α5β1 integrin antibody comprising:

 a heavy chain comprising an amino acid sequence having at least 95% identity to a sequence selected from the group consisting of SEQ ID NOs: 1, 16, 20, 25, 28, and 31; a light chain comprising an amino acid sequence having at least 95% identity to a sequence selected from the group consisting of SEQ ID NOs: 7, 18, 22, 26, and 32; and wherein the antibody inhibits angiogenesis stimulated by VEGF.
- 56. (Currently Amended) The anti-α5β1 integrin antibody of claim 55, wherein the light and heavy chain polypeptide sequences comprise amino acid sequences having at least 95% identity to SEQ ID NOs: 26 and 25, respectively.

- 57. (Currently Amended) The anti-α5β1 integrin antibody of claim 55, wherein the light and heavy chain polypeptide sequences comprise amino acid sequences having at least 95% identity to SEQ ID NOs: 28 and 26 and 28, respectively.
- 58. (Currently Amended) The anti-α5β1 integrin antibody of claim 55, wherein the light and heavy chain polypeptide sequences comprise amino acid sequences having at least 95% identity to SEQ ID NOs: 32 and 31, respectively.
- 59. (Currently Amended) A chimeric or humanized anti-α5β1 integrin antibody comprising: a heavy chain comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 20, 25, 28, and 31; and a light chain comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 22, 26, and 32.
- 60. (Previously Presented) The chimeric or humanized anti-α5β1 integrin antibody of claim 59, wherein the heavy chain polypeptide sequence comprises SEQ ID NO: 25 and the light chain polypeptide sequence comprises SEQ ID NO: 26.
- 61. (Previously Presented) The chimeric or humanized anti-α5β1 integrin antibody of claim 59, wherein the heavy chain polypeptide sequence comprises SEQ ID NO: 28 and the light chain polypeptide sequence comprises SEQ ID NO: 26.
- 62. (Previously Presented) The chimeric or humanized anti-α5β1 integrin antibody of claim 59, wherein the heavy chain polypeptide sequence comprises SEQ ID NO: 31 and the light chain polypeptide sequence comprises SEQ ID NO: 32.
- 63. (Previously Presented) A pharmaceutical composition comprising an anti-α5β1 integrin antibody according to any one of claims 51, 55, or 59, and a physiologically acceptable carrier.
- 64. (Withdrawn) A method for treating an ocular disease resulting in vascularization, said method comprising administering a dose of a pharmaceutical composition comprising an effective amount of an anti-α5β1 integrin antibody according to any one of claims 51, 55, or 59, and a physiologically acceptable carrier.

- 65. (Withdrawn) The method of claim 64, wherein the ocular disease is selected from the group consisting of macular degeneration, diabetic retinopathy, and choroidal neovascularization.
- 66. (Withdrawn) The method of claim 64, wherein the ocular disease is macular degeneration.
- 67. (Withdrawn) The method of claim 64, wherein the ocular disease is associated with secretion of VEGF.
- 68. (Withdrawn) The method of claim 64, wherein the treatment comprises intravenous injection.
- 69. (Withdrawn) The method of claim 64, wherein the treatment comprises intravitreal injection.
- 70. (Withdrawn) The method of claim 69, wherein the effective amount of anti- α 5 β 1 integrin antibody in each dose is at least about 100 μ g.
- 71. (Withdrawn) The method of claim 69, wherein the effective amount of anti- α 5 β 1 integrin antibody in each dose is at least about 300 μ g.
- 72. (Withdrawn) The method of claim 69, wherein the treatment comprises intravitreal injection into one eye, whereby the antibody contacts both eyes.
- 73. (Withdrawn) A nucleic acid encoding a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 2-6, 8-12, 16, 18, 20, 22, 25, 26, 28, 31, and 32.
- 74. (Withdrawn) A vector comprising at least one nucleic acid sequence selected from the group consisting of SEQ ID NOs: 15, 17, 19, 21, 23, 24, 27, 29, and 30.
- 75. (Withdrawn) The vector of claim 74, wherein the nucleic acid sequence comprises SEQ ID NOs: 23 and 24.
- 76. (Withdrawn) The vector of claim 74, wherein the nucleic acid sequence comprises SEQ ID NOs: 27 and 24.
- 77. (Withdrawn) The vector of claim 74, wherein the nucleic acid sequence comprises SEQ ID NOs: 29 and 30.

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78. (Withdrawn) A cell transformed by an expression vector comprising one or more of the nucleic acids comprising a sequence selected from the group consisting of SEQ ID NOs: 15, 17, 19, 21, 23, 24, 27, 29 and 30.